

Drug	Schedule
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Methadone (9250)	II
Methadone-intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II

DEA has considered the factors in 21 U.S.C. § 823(a) as well as information provided by other bulk manufacturers, and determined that the registration of Ganes Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 C.F.R. §§ 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: February 21, 1997.

Gene R. Haislip,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.

[FR Doc. 97-4895 Filed 2-26-97; 8:45 am]

BILLING CODE 4410-09-M

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 21, 1996, and published in the Federal Register on

November 14, 1996, (61 FR 58424), Norac Company, Inc., 405 S. Motor Avenue, Azusa, California 91702, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of tetrahydrocannabinols (7370), a basic class of controlled substance listed in Schedule I.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Norac, Company, Inc. to manufacturer tetrahydrocannabinols is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 C.F.R. §§ 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated February 6, 1997.

Gene R. Haislip,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.

[FR Doc. 97-4897 Filed 2-26-97; 8:45 am]

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Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 21, 1996, and published in the Federal Register on November 14, 1996, (61 FR 58424), Nycomed, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug

Enforcement Administration (DEA) to be registered as a bulk manufacturer of meperidine (9230), a basic class of controlled substance listed in Schedule II.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Nycomed, Inc. to manufacture meperidine is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 CFR §§ 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: February 7, 1997.

Gene R. Haislip,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.

[FR Doc. 97-4896 Filed 2-26-97; 8:45 am]

BILLING CODE 4410-09-M

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated July 31, 1996, and published in the Federal Register on August 8, 1996, (61 FR 41427), U.S. Drug Testing, Inc., 10410 Trademark Street, Rancho Cucamonga, California 91730, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Heroin (9200)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Phencyclidine (7471)	II
1-Piperidinocyclohexanecarbonitrile (8603)	II
Benzoylcegonine (9180)	II
Morphine (9300)	II

DEA has considered the factors in 21 U.S.C. 823(a), as well as information provided by other bulk manufacturers, and determined that the registration of U.S. Drug Testing, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 C.F.R. 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer

of the basic classes of controlled substances listed above is granted.

Gene R. Haislip,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.

[FR Doc. 97-4894 Filed 2-26-97; 8:45 am]

BILLING CODE 4410-09-M

DEPARTMENT OF LABOR

Employment Standards Administration

Proposed Collection; Comment Request

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and Federal agencies with an